APPENDIX F

Copy of August 21, 1992 Amendment from

Parent Application Serial No. 07/932,275

F-1 9116-335



P&G Case 4233C - Appleby et al.

Serial No.

Filed August 19, 1992

File Wrapper Continuation of USSN 07/580,706; Filed

Sept. 11, 1990

Received:
Amendment Requesting Interference Pursuant to

37 CFR §1.607, with 2 copies of transmittal sheet

Copy of U.S. 5,043,438 Certificate of Mailing

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IN THE UNITED STATES PATENT & TRADEMARK OFFICE RESPONSE/AMENDMENT

THE COMMISSIONER OF PATENTS AND TRADEMARKS Washington, D.C. 20231

Case Docket No. 4233C Parent in Group: 1803 Examiner: E. White

Dear Sir:

Transmitted herewith is an Amendment Requesting Interference Pursuant to 37 C.F.R. §1.607 for the File Wrapper Continuation patent application:

Inventor(s): DONALD B. APPLEBY et al.

Serial No.:

Date Filed: August 19, 1992

As a File Wrapper Continuation of Ser. No. 07/580,706; on

Filed September 11, 1990

Title: POLYOL POLYESTER SYNTHESIS

1. [] No additional fee is known to be required. 2. [X] The fee has been calculated as shown below: I hereby certify that this correspondence is being deposited with the United States Postal Service as first class mail in an envelope addressed to: Commissioner of Patents and Trademarks, Washington, D. C. 20231.

George W. Allen Name of Attorney Signature of Attorney

3/21/32 Date of Signature

OTUED THAN A

	(Col. 1)		(Col. 2)	(Col. 3)	SMALL ENTITY	
	CLAIMS REMAINING AFTER AMENDMENT		HIGHEST NO. PREVIOUSLY PAID FOR	PRESENT EXTRA*	RATE	FEE
TOTAL	* 56	MINUS	** 40	= 16	x \$ 20 =	\$320.00
INDEP.	* 3	MINUS	*** 3	= 0	x \$ 60 =	\$ 0
FIRST PRESENTATION OF MULTIPLE DEP. CLAIM					+ \$120 =	\$ 0
					TOTAL	\$320.00

If the entry in Col. 1 is less than the entry in Col. 2, write "O" in Col. 3.

** If the highest number of total claims previously paid for is less than 20, write "20" in this space.

*** If the highest number of independent claims previously paid for is less than 3, write "3" in this space.

The "Highest Number Previously Paid For" (Total or Independent) is the highest number found from the equivalent box in Col. 1 of a prior amendment or the number of claims originally filed.

- The Commissioner is hereby petitioned under 37 CFR §1.136(a) to grant any extension of time needed for timely response to the Office Action in the above-identified application to preserve pendency of said application. The processing fee under 37 CFR §1.17 has been determined as follows: \$_____ for a ____ month extension of
- The Commissioner is hereby authorized to charge payment of the following fees associated with this communication or credit any overpayment to Deposit Account No. 16-2480. A duplicate copy of this sheet is attached.

Any patent application processing fees under 37 CFR §1.16. Any patent application processing fees under 37 CFR §1.17. a. [x] b. [x]

The Commissioner is hereby authorized to make any additional copies of this sheet needed to accomplish the purposes provided for herein and to charge any fee for such copies to Deposit Account No. 16-2480.

George W. Allen

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Cincinnati, Ohio

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:

Donald B. Appleby et al.

Serial No.

: Group Art Unit 1803

Filed: August __, 1992

: Examiner: Mr. E. White

as a File Wrapper Continuation of

Serial No.: 07/580,706 Filed: September 11, 1990

For: POLYOL POLYESTER SYNTHESIS

AMENDMENT REQUESTING INTERFERENCE PURSUANT TO 37 C.F.R. §1.607

Hon. Commissioner of Patents and Trademarks Washington, D.C. 20231

Sir:

Please amend the above-identified file wrapper continuation application by adding the following claims for the purpose of interference with U.S. Patent No. 5,043,438, a copy of which is enclosed:

IN THE CLAIMS:

Please add the following new claims:

-- 63. A process for the synthesis of polyol fatty-acid polyesters by reacting a polyol and a fatty-acid lower-alkyl ester under substantially solvent free conditions in the presence of a catalyst and an emulsifier, the process comprising

an initial reaction stage which is carried out under such conditions that the reaction mixture in said initial stage is in steady-state, with continuous introduction of reactants comprising polyol and fatty-acid lower-alkyl ester, and continuous removal of products comprising reaction mixture having a degree of esterification of about 10% or more and volatile alcohol formed during the initial reaction stage, and

one or more subsequent reaction stages in which the reaction mixture from said initial stage is further reacted to said polyol fatty-acid polyesters. --

- -- 64. The process according to Claim 63 wherein the reaction mixture from the initial stage is further reacted to said polyol fatty acid polyesters after combining with any remaining part of the fatty-acid lower-alkyl ester reactant. --
- -- 65. A process according to Claim 63 wherein the emulisfier is an alkali metal soap. --
- -- 66. A process according to Claim 63 wherein the alkali metal soap is selected from the group of soaps having a chain length within the range of from 8 to 22 carbon atoms. --
- -- 67. A process according to Claim 63 wherein the fatty-acid lower-alkyl ester is a fatty-acid methyl ester. --
- -- 68. A process according to Claim 63 wherein the catalyst is selected from the group consisting of potassium hydroxide and carbonates of potassium and sodium. --

- -- 69. A process according to Claim 63 wherein the reaction mixture in said initial reaction stage has a degree of esterification of within the range of from 10 to 60%.
- -- 70. A process according to Claim 63 wherein the reaction mixture in said initial stage does not contain any substantial amount of solvent. --
- -- 71. A process according to Claim 63 wherein the reaction temperature in said initial stage is maintained at a level of within the range of from 130° to 140°C. --
- -- 72. A process according to Claim 63 wherein the average residence time of the reaction mixture in said initial stage is caused to be about 1.5 hours. --
- -- 73. A process according to Claim 63 wherein the molar ratio of catalyst to polyol in said initial reaction stage is within the range of from about 0.01:1 to about 0.5:1. --
- -- 74. A process according to Claim 63 wherein the molar ratio of emulsifier to polyol in said initial reaction stage is within the range of from 0.2:1 to 0.6:1. --
- -- 75. A process according to Claim 63 for the synthesis of polyol fatty-acid polyesters having a degree of esterification of at least about 70%. --
 - -- 76. A process according to Claim 63 wherein the polyol is sucrose. --
- -- 77. A process according to Claim 76 wherein the molar ratio of fatty-acid lower-alkyl ester to sucrose is within the range of from 7.2:1 to 15:1. --

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-- 78. A process according to Claim 63 wherein said initial reaction stage is fully separate from said one or more subsequent reaction stages. --

REMARKS

Upon entry of this amendment, Claims 1-6, 8-11, 13-15, 17-18, 22-24, 27-32, 40-45, 48-51, 54-58 and 62-78 are currently pending in the present file wrapper continuation application, including independent Claims 1, 62 and 63. Claim 62 was added by the First Preliminary Amendment. Claims 63-78 have been added by the present amendment.

New Claims 63-78 presented herein claim in whole or at least in part substantially the same subject matter as Claims 1-9, 13-15, and 17-20, respectively, of Buter U.S. Patent No. 5,043,438. Since the Buter '438 patent issued on August 27, 1991, the one-year time limit imposed by 35 U.S.C. §135(b) has been met.

Applicants point out that the Buter '438 patent is the subject of a recent Request for Reexamination, notice of which was published at 1139 OG 8 (June 2, 1992) (copy enclosed). See MPEP §§2307.04, 2284. Applicants' counsel have reviewed this Request for Reexamination on file in the PTO, and are not aware of any references more pertinent than those of record in the Request for Reexamination and in applicants' parent application, Serial No. 07/580,706.

Support for new Claims 63-78 can be found in the specification of the present application as shown in the following table:

APPLICATION CLAIMS

SUPPORT IN APPLEBY ET AL. APPLICATION

63. A process for the synthesis of polyol fatty-acid polyesters

Page 1, lines 13-14: "This invention relates to improved synthesis of higher polyol fatty acid polyesters."

SUPPORT IN APPLEBY ET AL. APPLICATION

by reacting a polyol and

Page 2, lines 29-31: "The present invention relates to . . . interesterifying (a) polyol containing more than about four esterifiable hydroxy groups. . . "

a fatty-acid lower-alkyl ester

Page 2, lines 32-33: "... and (b) fatty acid ester of easily removable alcohol."

Page 5, lines 26-33: "As used herein, the terms 'fatty acid ester(s)' and 'ester reactant(s)' . . . are preferably esters of volatile alcohols, e.g., the C₁-C₄ (preferably methyl) [i.e., lower alkyl] . . . esters of fatty acids containing about eight or more carbon atoms."

under substantially solvent free conditions

Page 1, lines 13-16: "This invention relates to improved synthesis of higher polyol fatty acid polyesters . . . via transesterification reactions that do not use a solvent to form a homogeneous reaction mix."

Page 2, lines 34-35: "in the absence of any substantial amount of unreactive solvent."

in the presence of a catalyst and

Page 3, lines 5-7: "The process is a continuous process in which the initial catalyst level is from about 0.01 to about 0.5 moles of catalyst per mole of polyol. . . . "

Page 11, lines 25-27: "The basic catalysts generally suitable for use in preparing the polyol polyesters described herein are

Page 3, lines 9-10: "The initial level of

an emulsifier,

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the process comprising: an initial reaction stage

which is carried out under such conditions that the reaction mixture in said initial stage is in steady-state with

continuous introduction of reactants

soap emulsifier in the first stage of the reaction is from about 0.001 to about 0.6, preferably from about 0.2 to about 0.4 moles per mole of polyol."

Page 24, lines 37-38: "In general it is desirable, and even preferred, to effect the reaction in at least two stages."

Page 19, lines 19-20: "It is desirable in the initial stage, or stages, of the process, and especially of a continuous process, to have a relatively high degree of completion."

Page 1, lines 13-17: "This invention relates . . . more specifically to a continuous process. . . ."

Page 36, lines 5-6: "The continuous reactor is started up and allowed to reach steady state."

Page 42, lines 9-14: "... continous operation is started with the feed slurry introduced into the first reactor and the intermediate product from the first reactor cascading to the second and so on down the train. The reactors are allowed to reach steady-state with the average composition at steady-state exiting from each reactor about the following. . . ."

Page 19, lines 14-18: "In a continuous reaction, the individual reactants can be added to the first stage at a rate that maintains the desired degree of esterification and yet provides sufficient yield from the first stage to maintain the

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reaction in the subsequent stage, or stages."

Page 20, lines 5-7: "... reactants are continually added and the product is removed at rates that maintain the desired level of esterification."

comprising polyol and fatty-acid lower-alkyl ester, and

Page 24, lines 27-31: "an initial heterogenous reaction mixture comprises . . . polyol [and] . . . fatty acid ester "

Page 5, lines 26-31: "... the terms 'fatty acid ester(s)' ... are preferably esters of volatile alcohols, e.g., the C_1 - C_4 (preferably methyl) ... esters of fatty acids. ..."

continuous removal of products comprising reaction mixture having a degree of esterification of 10% or more and

Page 42, lines 9-14: "... continous operation is started with the feed slurry introduced into the first reactor and the intermediate product from the first reactor cascading to the second and so on down the train. The reactors are allowed to reach steady-state with the average composition at steady-state exiting from each reactor about the following..."

Page 20, lines 5-7: "... reactants are continually added and the product is removed at rates that maintain the desired level of esterification."

Page 19, lines 3-6: "It is highly desirable to conduct the initial stage... to maintain the degree of esterification between about 10% (preferably 20%) and about 70%...."

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Page 36, lines 5-11: "The continuous reactor is started up and allowed to reach steady state. . . . At steady state the product exiting from the continuous reactor has an average composition of about 3.05 fatty acid chains per molecule of sucrose. . . ." "Degree of esterification" is defined at page 2, lines 18-23, as the percentage "of the total number of available hydroxy groups on the polyol [that] are esterified with a fatty acyl radical." Thus, 3.05 fatty acid chains per molecule of sucrose (which has 8 available hydroxyl groups) is equivalent to 3.05 ÷ 8, or 38.125% esterification.

See also Examples 14 (pages 41-42) and 16 (pages 43-44), giving details of continuous processes and steady state composition of the effluent partial esters.

Page 3, line 37 - Page 4, line 5: "The said easily removable alcohol is a volatile alcohol . . . and the removal of the volatile alcohol produced by the reaction between the ester reactant and the polyol is assisted by increasing the mass transfer area of the reaction mixture . . . and/or sparging. . . . "

Page 24, lines 37-38: "In general it is desirable, and even preferred, to effect the reaction in at least two stages. . . ."

Page 19, lines 14-18: "In a continuous reaction, the individual reactants can be added to the first stage at a rate that maintains the desired degree of esterification and yet provides sufficient

volatile alcohol formed during the initial reaction stage, and

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one or more subsequent reaction stages in which the reaction mixture from said initial stage is further reacted to said polyol fatty-acid polyesters.

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yield from the first stage to maintain the reaction in the subsequent stage, or stages."

Page 4, lines 23-28: "The final stage, or stages, of the reaction are carried out . . . after the degree of esterification of said polyol has reached at least about 50% to achieve a final degree of esterification of at least about 70%, preferably at least about 95%."

Page 21, lines 25-37: "The final stage, or stages, of the reaction should be carried out under . . . conditions to . . . achieve high degrees of esterification. . . . The final degree of esterification should be at least about 70%, preferably at least about 98%."

Page 19, lines 31-36: "Maintenance of the appropriate composition for solubilizing the polyol is assisted by withholding a portion of the ester reactant from this initial stage In the initial stage, it is preferable to use only about 30% to about 70% of the total ester reactant, with the remainder being added in the later stages."

Pages 25, lines 1-3: "In any later stage, additional fatty acid esters . . . can be added."

65. A process according to Claim 63 wherein the emulsifier is an alkali metal soap.

64. The process according to Claim 63

combining with any remaining part of the

wherein the reaction mixture from the

initial stage is further reacted to said

fatty-acid lower-alkyl ester reactant.

polyol fatty-acid polyesters after

Page 11, lines 7-8: "Alkali metal soaps are typically, and preferably, used as emulsifiers in the processes described herein."

66. A process according to Claim 65 wherein the alkali metal soap is selected from the group of soaps having a chain length within the range of from 8 to 22 carbon atoms.

67. A process according to Claim 63 wherein the fatty-acid lower-alkyl ester is a fatty-acid methyl ester.

68. A process according to Claim 63 wherein the catalyst is selected from the group consisting of potassium hydroxide and carbonates of potassium and sodium.

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69. A process according to Claim 63 wherein the reaction mixture in said initial reaction stage has a degree of esterification of within the range of from 10 to 60%.

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- Page 11, lines 9-12: "the term 'alkali metal fatty acid soap' is intended to include the alkali metal salts of saturated and unsaturated fatty acids having from about 8 to about 18 carbon atoms."
- Page 11, lines 20-24: "In a preferred process, . . . it is highly desirable that any soap present be an alkali metal . . . salt of hydrogenated fatty acids containing from about 16 to about 22 carbon atoms."
- Page 5, lines 26-34: "As used herein, the terms 'fatty acid ester(s)' and 'ester reactant(s)' are intended to include any compound wherein the alcohol portion is easily removed, . . . but are preferably esters of volatile alcohols, e.g., the C₁-C₄ (preferably methyl) . . . esters of fatty acids. . . . Methyl esters are the most highly preferred ester reactants."
- Page 11, lines 35-38: "In another particularly preferred embodiment of the present invention, the basic catalyst used in the reaction is potassium carbonate, sodium carbonate. . . ."
- Page 34, lines 32-33: "The catalyst in this reaction is the residual KOH [potassium hydroxide] in the soap."
- Page 19, lines 3-6: "It is highly desirable to conduct the initial stage . . . to maintain the degree of esterification between about 10% (preferably 20%) and about 70%, preferably between about 35% and about 60%."

70. A process according to Claim 63 wherein the reaction mixture in said initial stage does not contain any substantial amount of solvent.

- 71. A process according to Claim 63 wherein the reaction temperature in said initial stage is maintained at a level of within the range of from 130° to 140°C.
- 72. A process according to Claim 63 wherein the average residence time of the reaction mixture in said initial stage is caused to be about 1.5 hours.

- 73. A process according to Claim 63 wherein the molar ratio of catalyst to polyol in said initial reaction stage is within the range of from about 0.01:1 to about 0.5:1.
- 74. A process according to Claim 63 wherein the molar ratio of emulsifier to polyol in said initial reaction stage is within the range of from 0.2:1 to 0.6:1.

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- Page 2, lines 34-35: "in the absence of any substantial amount of unreactive solvent."
- Page 1, lines 13-16: "This invention relates to improved synthesis of higher polyol fatty acid polyesters . . . that do not use a solvent to form a homogenous reaction mix."
- Page 18, lines 2-5: "It is highly preferable to run the reaction in the initial stages at temperatures between about 265°F (130°C) and about 285°F (140°C), preferably between about 270°F (132°C) and about 275°F (135°C)."
- Page 35, lines 30-32: "Examples 9 and 10 make use of a 3-liter continuous stirred tank reactor (CSTR) to perform the first stage of the reaction. The residence time of this reactor is about 1.5 hours."
- Page 36, lines 6-7: "The reactor conditions are . . . about 1.5 hours residence time."
- Page 3, lines 5-7: "The process is a continuous process in which the initial catalyst level is from about 0.01 to about 0.5 moles of catalyst per mole of polyol, preferably from about 0.01 to about 0.1 mole of catalyst per mole of polyol."
- Page 11, lines 7-8: "Alkali metal soaps are typically, and preferably, used as emulsifiers in the processes described herein."

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- 75. A process according to Claim 63 for the synthesis of polyol fatty-acid polyesters having a degree of esterification of at least about 70%.
- 76. A process according to Claim 63 wherein the polyol is sucrose.
- 77. A process according to Claim 76 wherein the molar ratio of fatty-acid lower-alkyl ester to sucrose is within the range of from 7.2:1 to 15:1.

Page 15, lines 15-17: "The level of soap in the first stage of the reaction is desirably from about 0.001 to about 0.6, preferably from about 0.2 to about 0.4 moles of soap per mole of polyol."

Page 21, lines 36-37: "The final degree of esterification should be at least about 70%, preferably at least about 98%."

Page 4, lines 34-35: "In practicing the process disclosed herein, sucrose is the most highly preferred polyol."

Page 25, lines 3-7: "In any second, or later step, additional fatty acid ester can be added to raise the overall ratio of fatty acyl groups to the esterifiable hydroxy sites on the polyol to from about 0.9:1 to about 1.4:1 [since sucrose has 8 esterifiable sites, this translates to about 7.2:1 to about 11.2:1 moles ester to moles sucrose], preferably from about 1:1 to about 1.2:1 [about 8:1 to about 9.6:1 moles ester to moles sucrose]."

Page 30, lines 17-28: "To perform the first stage of the reaction, about 148 grams of partially hardened soybean methyl esters (0.5 mole) are added to the reactor To this mixture is added about 34.2 grams of crystalline sucrose (0.1 mole). . . . To perform the second stage of the reaction, about 207 grams of partially hardened soybean methyl esters (0.7 mole) . . . are added to the reactor." [Total esters 1.2 mole, total sucrose 0.1 mole; overall molar ratio = 12:1.]

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78. A process according to Claim 63 wherein said initial reaction stage is fully separate from said one or more subsequent reaction stages.

Page 34, lines 34-37: "About . . . 25 grams of sucrose (0.073 moles) . . . and about 320 grams of partially hardened soybean methyl esters (1.094 moles) are added to the reactor." [Molar ratio = 15:1].

Page 24, lines 37-38: "In general it is desirable, and even preferred, to effect the reaction in at least two stages."

Page 21, lines 13-21: "It is advantageous ... to carry out the reaction in a series of at least two, preferably from two to about eight, more preferably from three to about eight, reaction vessels. ... Separate vessel/reactors allow for closer control. ... Separate vessels are also particularly desirable. ..."

Page 40, lines 4-8: "This example demonstrates the ability to utilize a series of three Continuously Stirred Tank Reactors (CSTR) for the first stage of a reaction to make sucrose polyesters . . . and then completing the second stage of the reaction in a batch reactor."

Pursuant to the provisions of 37 C.F.R. §1.607, and in view of the foregoing amendments and accompanying remarks, Applicants respectfully request that an interference be declared between the common claimed subject matter of the present application and Buter U.S. Patent No. 5,043,438.

In particular, it is proposed that the interference be declared with a single count, reading substantially as Buter '438 patent Claim 1 and applicants' Claim 63, as follows:

PROPOSED COUNT I

A process for the synthesis of polyol fatty-acid esters by reacting a polyol and a fatty-acid lower-alkyl ester under substantially solvent free conditions in the presence of a catalyst and an emulsifier, the process comprising

an initial reaction stage which is carried out under such conditions that the reaction mixture in said initial stage is in steady-state, with continuous introduction of reactants comprising polyol and fatty-acid lower-alkyl ester, and continuous removal of products comprising reaction mixture having a degree of esterification of about 1% or more and volatile alcohol formed during the initial reaction stage of the synthesis, and

one or more subsequent reaction stages in which the reaction mixture from said initial stage is further reacted to said polyol fatty-acid esters.

It is proposed that the interference be declared with Buter '438 patent Claims 1-21 and so much of applicants' Claims 63-78 as correspond exactly to the Buter claims be designated as unpatentable over the interference count.

Proposed Count I is written in the language of applicants' Claim 63, but is believed broad enough to encompass both applicants' Claim 63 and Buter Claim 1. Like Buter Claim 1, the count recites a process for the synthesis of "esters," rather than being limited to the subcategory "polyesters," and recites a degree of esterification in the first stage of the reaction of "about 1% or more," rather than being limited to applicants' narrower

range "about 10% or more." Thus, the proposed count is not narrower than Buter Claim 1, the broadest claim of the Buter patent. See 37 C.F.R. §1.606.

The differences in language between applicants' Claim 63 and Buter patent

Claim 1 are such that both of these claims are nevertheless drawn to the same subject matter.

The language of Claim 63 of the present application is identical to that of Buter '438 patent Claim 1, except that:

- (1) Claim 63 is directed to a process for the synthesis of --polyol fatty-acid polyesters-- instead of "polyol fatty-acid esters". Both claims are drawn to substantially the same subject matter, however, since the process claimed in the Buter '438 patent can be used to synthesize both "esters" and "polyesters", the only difference being the degree of completion to which the reaction is allowed to run. Indeed, the Buter patent is directed to the synthesis of both polyol fatty acid esters and polyesters; Buter dependent Claim 16 specifically claims the synthesis of polyesters. Moreover, the Buter patent indicates that the process is particularly directed to the synthesis of polyesters, although it is suitable also for synthesis of oligoesters. Col. 6, Ins. 53-56. Buter patent Example 1 discloses the synthesis of sucrose polyester with 96.6% conversion. Col. 8, Ins. 18-20.
- (2) Claim 63 (as well as Claims 68 and 73) omits the word "transesterification" before "catalyst." This is not a difference in substance, since the reaction which is described in the instant application, in which the catalyst used, is in fact a transesterification reaction, as indicated on page 1, lines 13-15. One of ordinary skill in the art would know that the type of catalyst called for in a transesterification reaction is inherently a "transesterification catalyst."

- (3) Claim 63, as well as Claims 64, 69-74 and 78, refer to initial and subsequent reaction "stages," like Buter Claim 1, but omit the terminology referring to first and subsequent reaction "zones." This is simply a difference in wording which does not change the fact that substantially the same subject matter is involved. Both applicants' disclosure and the Buter '438 patent are directed to a reaction having at least two distinct stages. (See discussion of support for Claim 63, above.) The Buter patent states that the first stage is carried out in a "first reaction zone" which may be a reaction vessel separate from one or more further reaction vessels or may be part of multi-zone continuous esterification equipment which, in turn, may be "a serial sequence of separate reaction vessels." Col. 3, lines 3-8. Similarly, the instant application discloses that a "series of reaction vessels and/or separate reactors" should be used in the reaction (Page 21, lines 16-17) and that in one embodiment, the reaction is done "using two backmix reactors in series for the initial stage."
- -continuous removal of products-- rather than "in-going reactant streams" and "out-going product streams." This difference again merely represents two ways of of describing the same aspect of the same reaction scheme. Claim 63 uses the term --continuous-- instead of Buter's term "massbalanced" to refer to the fact that the first stage of the reaction involves a continuous inflow of reactants and a continuous outflow of the steady state reaction mixture of product and reactants. This does not represent a change in substance. A person of ordinary skill in the art would know that for a process to be in steady state, the rate of mass continuously entering the system must equal the rate of mass continuously leaving, i.e. there

must inherently be "mass balance". See, e.g., Unit Operations of Chemical Engineering, 3d Ed. by McCabe and Smith, (McGraw-Hill, 1976) at page 66 (copy enclosed). A continuous process must operate in steady state and be mass balanced except during start-up. See, e.g., Elementary Principles of Chemical Processes by Felder et al. (Wiley 1978), page 82 (copy enclosed). Felder et al. define a "continuous process" as a process where "[t]he inputs and outputs flow continuously through the duration of the process." Felder et al. indicate that if "all the variables [including rate of inflow and outflow] in a process do not change with time, except possibly for minor fluctuations about constant mean values, the process is said to be operating at steady state." Felder et al. state that "[c]ontinuous processes are usually run as close to steady state as possible; unsteady-state (transient) conditions exist during the start-up of a process and following changes - intentional or otherwise - in process operating conditions." Thus, a person of ordinary skill in the art would know that where there is "continuous introduction of reactants" and "continuous removal of products" and the reaction is in steady state (i.e., after start-up), the inflows and outflows must inherently be "massbalanced." Otherwise, the reaction vessel would either fill up and reach its capacity, or empty out, and the process would cease to be continuous or in steady state. The Buter '438 patent at col. 2, lns. 24 and 42 makes clear that the patent is directed to a continuous process, as is the instant application (page 1, line 17). Indeed, in the prosecution history of the Buter '438 patent, the Examiner noted in his "reasons for allowance" that Buter claimed "a continuous initial reaction stage."

(5) Claim 63 recites a --a degree of esterification of about 10% or more-instead of "a polyol conversion of 1% or more" as set out in Buter Claim 1. "Polyol

conversion" is defined in the Buter patent as "the percentage of polyol hydroxyl groups of the original polyol that on average have been esterified with fatty acids." Buter patent col.

1, In. 32-37. Similarly, at page 2, lines 18-23 of the present application, the term "degree of esterification" is used to mean the percentage "of the total number of available hydroxy groups on the polyol [that] are esterified with a fatty acyl radical." Applicants' specification does not disclose polyol conversion below 10% in the initial stage. But even the Buter patent makes clear that the 1% is not a critical lower limit. Indeed, Buter teaches that the preferred first stage conversion is at least 10% (col. 3, line 31). Moreover, the lowest polyol conversion disclosed in the examples of the Buter patent for the first stage of the process is 10.3% (Example 4, col. 9, ln. 56), while other Buter examples show first stage conversions of 25.6% (Example 2), 32.2% (Example 3), and 41.5% (Example 5). Under these circumstances, an interference is properly declared between applicants' Claim 63 and Buter's Claim 1 using the 1% value set out in Buter's claim. See MPEP §2309.01.

(6) Claim 63 refers to the --volatile alcohol-- formed during the initial stage of the synthesis instead of a "lower-alkyl alcohol." Given that the reactants are a polyol and a lower-alkyl fatty acid ester, the products of the transesterification reaction will inherently be polyol fatty-acid esters and a lower-alkyl alcohol, which is inherently volatile. Moreover, while Buter Claim 1 refers to the lower-alkyl alcohol as part of the out-going product "streams," the Buter '438 patent makes clear that it is removed as a vapor. '438 patent col. 5, ln. 55 - col. 6, ln. 7.

The Dependent Claims

The language of Claim 64 is identical to that of Buter '438 patent Claim 2 except that it recites polyol fatty-acid --polyesters-- instead of "esters," to agree with Claim 63, and it recites the phrase --the fatty-acid lower-alkyl ester reactant-- instead of "reactants". This is not a change in substance, since when the Buter patent discloses the addition of reactants in the later stages, it specifies "in particular the fatty-acid lower-alkyl ester." Col. 3, Ins. 53-57.

The language of Claim 66 is identical to that of Buter '438 patent Claim 4

except that it recites a partially overlapping range, i.e. from --8 to 22-- carbon atoms instead

of "6 to 14" carbon atoms, and omits the phrase "short chain" preceding the word "soaps."

The language of Claim 68 is identical to that of Buter '438 patent Claim 6

except that it omits the word "transesterification" before "catalyst", as discussed above, and recites --potassium hydroxide and carbonates of potassium and sodium-- instead of "hydroxides, carbonates and bicarbonates of potassium and sodium."

The language of Claim 69 is identical to that of Buter '438 patent Claim 7 except that it recites a range of --10 to 60%--, which encompasses Buter's narrower range of "10 to 40%". In addition, Claim 69 uses the terminology --stage-- instead of "zone" and -- degree of esterification-- instead of "degree of polyol conversion." As discussed above with respect to Claim 63, this terminology is equivalent.

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The language of Claim 70 is identical to that of Buter '438 patent Claim 8 except that it uses --stage-- instead of "zone" as discussed above, and includes the phrase

--does not contain any substantial amount of solvent-- instead of "has a solvent level of 0.1% by weight or less." The Buter patent makes clear that its objective is to carry out the reaction "under substantially solvent-free conditions." Col. 1, lns. 7-8.

The language of Claim 71 is identical to that of Buter '438 patent Claim 9 except that it uses --stage-- instead of "zone" as discussed above, and recites a narrower temperature range of --130°C to 140°C--, within Buter's "120°C to 150°C."

The language of Claim 72 is identical to that of Buter '438 patent Claim 13 except that it uses --stage-- instead of "zone" as discussed above, and recites an average residence time of --about 1.5 hours--, within Buter's "range of from 1.5 to 2.5 hours."

The language of Claim 73 is identical to that of Buter '438 patent Claim 14

except that it uses --stage-- instead of "zone" and omits the word "transesterification" before catalyst, as discussed above, and recites a molar ratio range of --about 0.01:1 to about 0.5:1--, which encompasses Buter's narrower range of "0.1:1 to 0.3:1."

The language of Claim 74 is identical to that of Buter '438 patent Claim 15

except that it uses --stage-- instead of "zone" as discussed above, and recites a molar ratio of

--0.2:1 to 0.6:1--, within the broader range of "0.2:1 to 0.8:1" claimed by Buter.

The language of Claim 75 is identical to that of Buter '438 patent Claim 17 except that it recites a degree of esterification of --at least about 70%--, which encompasses Buter's narrower range of "90% or more". In addition, the claim's dependency has been changed to depend on Claim 63. In the Buter '438 patent, Claim 17 depends on Claim 16, which has no corresponding claim in the present application because it would have been redundant. Finally, Claim 75 uses the terminology --degree of esterification-- instead of

"polyol conversion." As discussed above with respect to Claim 63, these terms are

equivalent.

The language of Claim 77 is identical to that of Buter '438 patent Claim 19

except that it recites a molar ratio of --7.2:1 to 15:1--, partially overlapping Buter's range of

"10.5:1 to 18:1."

The language of Claim 78 is identical to that of Buter '438 patent Claim 20

except that it uses the terms --stage-- and --stages-- instead of "zone" and "zones" to describe

the first reaction stage and separate subsequent reaction stages, respectively, as discussed

with respect to Claim 63 above.

Finally, other than their numbering, and other than the fact that the claim on

which they depend, Claim 63, differs from Buter '438 patent Claim 1 as noted above, Claims

65, 67 and 76 correspond exactly to Claims 3, 5 and 18 of the Buter '438 patent.

It is therefore respectfully requested that an interference be declared between

the instant application and Buter U.S. Patent No. 5,043,438.

Respectfully submitted,

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Cincinnati, Ohio

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BASIC EQUATIONS OF FLUID FLOW

The principles of physics most useful in the applications of fluid mechanics are the mass-balance, or continuity, equations; the linear- and angular-momentum-balance equations; and the mechanical-energy balance. They may be written in differential form, showing conditions at a point within a volume element of fluid, or in integrated form applicable to a finite volume or mass of fluid. Only the integral equations are discussed in this chapter.

Certain simple forms of the differential equations of fluid flow are given in later chapters. A complete treatment requires vector and tensor equations outside the scope of the text. They underlie more advanced study of the phenomena, and they are discussed in many texts dealing with applied mechanics and transport processes. 1-3

Mass balance In steady flow the mass balance is particularly simple. The rate of mass entering the flow system equals that leaving, as mass can be neither accumulated nor depleted within a flow system under steady conditions.

STREAMLINES AND STREAM TUBES Discussions of fluid-flow phenomena are facilitated by visualizing, in the stream of fluid, fluid paths called *streamlines*. A streamline is an imaginary curve in a mass of flowing fluid so drawn that at every point on the curve

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